

Amendments to the Claims

1-38. (Cancelled)

39. (Currently amended) An oligonucleotide probe comprising a sequence of at least 10 contiguous nucleotides of a human mitochondrial genome, wherein the oligonucleotide probe comprises a ~~mutation selected from the group consisting of:~~ a mutation selected from the group consisting of: T → C at nucleotide 114; ΔC mutation at nucleotide 302; C → A at nucleotide 386; insert T at nucleotide 16189; A → C at nucleotide 16265; A → T at nucleotide 16532; C → T at nucleotide 150; T → C at nucleotide 195; ΔC at nucleotide 302; C → A at nucleotide 16182; C → T at nucleotide 16187; T → C at nucleotide 16519; G → A at nucleotide 16380; G → A at nucleotide 75; insert C at nucleotide 302; insert C → G at nucleotide 514; T → C at nucleotide 16172; C → T at nucleotide 16292; A → G at nucleotide 16300; A → G at nucleotide 10792; C → T at nucleotide 10793; C → T at nucleotide 10822; A → G at nucleotide 10978; A → G at nucleotide 11065; G → A at nucleotide 11518; C → T at nucleotide 12049; T → C at nucleotide 10966; G → A at nucleotide 11150; G → A at nucleotide 2056; T → C at nucleotide 2445; T → C at nucleotide 2664; T → C at nucleotide 10071; T → C at nucleotide 10321; T → C at nucleotide 12519; Δ 7 amino acids at nucleotide 15642; G → A at nucleotide 5521; G → A at nucleotide 12345; G → A at nucleotide 3054; T → C substitution at position 710; T → C substitution at position 1738; T → C substitution at position 3308; G → A substitution at position 8009; G → A substitution at position 14985; T → C substitution at position 15572; G → A substitution at position 9949; T → C substitution at position 10563; G → A substitution at position 6264; A insertion at position 12418; T → C substitution at position 1967; and T → A substitution at position 2290.

40. (Currently amended) An oligonucleotide primer comprising a sequence of at least 10 contiguous nucleotides of a human mitochondrial genome, wherein the oligonucleotide primer comprises a ~~mutation selected from the group consisting of:~~ a mutation selected from the group consisting of: T → C at nucleotide 114; ΔC mutation at nucleotide 302; C → A at nucleotide 386; insert T at nucleotide 16189; A → C at nucleotide 16265; A → T at nucleotide 16532; C → T at nucleotide 150; T → C at nucleotide 195; ΔC at nucleotide 302; C → A at nucleotide 16182; C → T at nucleotide 16187; T → C at nucleotide 16519; G → A at nucleotide 16380; GA at nucleotide 75; insert C at nucleotide 302; insert C → G at nucleotide 514; T → C at nucleotide 16172; C → T at

~~nucleotide 16292; A → G at nucleotide 16300; A → G at nucleotide 10792; C → T at nucleotide 10793; C → T at nucleotide 10822; A → G at nucleotide 10978; A → G at nucleotide 11065; G → A at nucleotide 11518; C → T at nucleotide 12049; T → C at nucleotide 10966; G → A at nucleotide 11150; G → A at nucleotide 2056; T → C at nucleotide 2445; T → C at nucleotide 2664; T → C at nucleotide 10071; T → C at nucleotide 10321; T → C at nucleotide 12519; A 7 amino acids at nucleotide 15642; G → A at nucleotide 5521; G → A at nucleotide 12345; G → A at nucleotide 3054; T → C substitution at position 710; T → C substitution at position 1738; T → C substitution at position 3308; G → A substitution at position 8009; G → A substitution at position 14985; T → C substitution at position 15572; G → A substitution at position 9949; T → C substitution at position 10563; G → A substitution at position 6264; A insertion at position 12418; T → C substitution at position 1967; and T → A substitution at position 2299.~~

41-117. (Cancelled)

118. (New) The oligonucleotide probe of claim 39 or primer of claim 40 which comprises a sequence of at least 12 contiguous nucleotides of a human mitochondrial genome.

119. (New) The oligonucleotide probe of claim 39 or primer of claim 40 which comprises a sequence of at least 14 contiguous nucleotides of a human mitochondrial genome.

120. (New) The oligonucleotide probe of claim 39 or primer of claim 40 which comprises a sequence of at least 16 contiguous nucleotides of a human mitochondrial genome.

121. (New) The oligonucleotide probe of claim 39 or primer of claim 40 which comprises a sequence of at least 18 contiguous nucleotides of a human mitochondrial genome.

122. (New) The oligonucleotide probe of claim 39 or primer of claim 40 which comprises a sequence of at least 20 contiguous nucleotides of a human mitochondrial genome.

123. (New) The oligonucleotide probe of claim 39 or primer of claim 40 which comprises a sequence of at least 22 contiguous nucleotides of a human mitochondrial genome.

124. (New) The oligonucleotide probe of claim 39 or primer of claim 40 which comprises a sequence of at least 24 contiguous nucleotides of a human mitochondrial genome.

125. (New) The oligonucleotide probe of claim 39 or primer of claim 40 which comprises a sequence of at least 26 contiguous nucleotides of a human mitochondrial genome.

126. (New) The oligonucleotide probe of claim 39 or primer of claim 40 which comprises a sequence of at least 30 contiguous nucleotides of a human mitochondrial genome.